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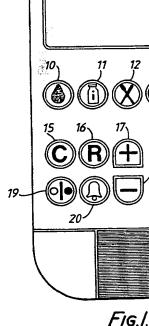
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Diabetes management system and apparatus.

A system and apparatus for efficient medical control of a medical condition such as diabetes comprises a recorder, an interface and a master computer. The master computer develops a programme of therapy which is downloaded into the recorder which then reminds the patient of any therapy due and records that the therapy has been effected. The record from the recorder is subsequently fed back to the master computer to improve or alter the therapy programme.



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DIABETES MANAGEMENT SYSTEM AND APPARATUS

This invention is concerned with apparatus for use in a system of outpatient management, i.e. a system for gathering, processing and analysing data to provide for an outpatient an individually tailored programme of treatment or medication.

The system described herein has been specifically designed for the control of diabetes but could be applied to many other conditions requiring outpatient monitoring.

In managing diabetes it is normal to derive for each patient the optimum programme of eating, insulin doses and exercise, the patient then following that programme in the course of their normal daily life and compensating for any departures from the normal programme. There are two major problems inherent in this, namely that the optimum programme is subject to change because of changing age or health of the patient or because of a change in the patient's daily routine, and that any compensation by the patient for a departure from the programme is at best an educated guess.

It is an object of the present invention to provide apparatus in the context of an overall management system that obviates or mitigates these problems.

The present invention is a system of outpatient management comprising a computer into which are loaded details of the patient, or his condition, a monitor in the possession of the patient and an interface, enabling the computer and the monitor to exchange information, the computer being programmed to produce from the loaded patient details a course of treatment or medication which is recorded in the patient's monitor, the monitor including a computer, a display and a keyboard and being programmed to use the recorded course to display at appropriate times the desired treatment or medication and to accept and record information entered by the patient on the keyboard.

The present invention is also a monitor for use by a patient in a system of outpatient management, the monitor comprising a computer, a keyboard and a display, the computer being adapted to store a course of medication or treatment, to display at appropriate times the desired medication or treatment and to accept and store keyboard input from the patient relating to the medication or treatment.

The present invention is further apparatus for measuring colour change comprising means for supporting an article subject to colour change, a light source and means for directing light from the source onto the article as a well defined spot, and means for receiving reflected light from the light source and measuring the intensity of the light.

An embodiment of the present invention will now be described by way of example, with reference to the accompanying drawings, in which:

Fig. 1 shows the face of a recorder used by a patient;

Fig. 2 shows the full display of the recorder of Fig. 1;

Fig. 3 is a block schematic diagram of the recorder of Fig. 1;

Fig. 4 is a cross sectional view of a blood glucose strip reader;

Fig. 5 is a plan view of a strip carrier used in the reader of Fig. 4;

Fig. 6 is a cross section on the line 6-6 of Fig. 5;

Fig. 7 is an exploded view of a reflection compartment used in the reader of Fig. 4;

Fig. 8 is a circuit diagram of the blood glucose strip reader; and

Fig. 9 is a block circuit diagram of an interface unit used with the recorder of Fig. 1.

The overall system of diabetes management comprises three elements, namely a patient operated monitor/recorder (hereinafter referred to simply as a recorder), a master computer for developing the optimum programme on the basis of the information recorded by the patient, and an interface unit used to transfer data between the recorder and the master computer or a printer or even a modem.

The master computer is simply a personal computer having a special programme and which is operated by the doctor or medical specialist to develop on the basis of the patient's data the optimum programme for that patient and to insert appropriate instructions into the patient's recorder.

The recorder itself, when the appropriate instructions have been entered from the master computer, prompts the patient to perform the actions according to the programme at the appropriate time. The patient also enters into the memory of the recorder information relating to insulin types and doses, diet, exercise, urine test results, hypoglycemic reactions and special events. The recorder also incorporates a blood glucose test strip reader and automatically stores the measured values in the memory. All the entries in memory are coded and labelled with date and time.

The keyboard of the recorder is illustrated in Fig. 1, the various symbols on the various keys 10 to 21 having the following meaning:-

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10 - Blood Glucose: Starts a procedure enabling read out of BG test strips according to manufacturer's procedure 11 - Insulin: Starts a procedure enabling review and/or entry of insulin guidelines 12 - Diet : Starts a procedure enabling review and/or entry of diet guidelines 13 - Exercise : Starts a procedure enabling review and/or entry of exercise guidelines 5 14 - Urine Test: Starts a procedure enabling entry of urine test results according to existing methods of urine testing 15 - C key: Starts a procedure enabling clearing or zero setting of the values in the display, preventing memory storage of displayed values, and enabling clock/calender setting 16 - R key: Starts a procedure enabling restoration of preset values, review of previous entered BG 10 values, and recall of previous preset guidelines 17 - + key: Enables the increase of values displayed and the recording of special events 18 - - key: Enables the decrease of values displayed and recording of hypoglycemic reactions 19 - on/off key: Enables on/off switching of the Romeo 20 - bell key: Switches alarm on or off 15 21 - M key: Starts procedure enabling to store the various displayed data into Romeo's memory. The monitor/recorder also has an LCD as shown in Fig. 2, the various symbols having the following meaning:-20 = Blood Glucose function active = Insulin function active 25 = Diet function active = Exercise function active 30 = Urine test function active И Ω = Alarm enabled PM/AM = Time indicators 35 = Memory full up indicators = Appointment time/Consult physician indicator 3 = Procent indicator 40 = Alpha Numeric digit = Numeric digit 45 = Seconds indicator = Floating point

In Fig. 3 is shown a block schematic diagram of the recorder.

BAT

M

R

= Battery Iow indicator

= M key operation

= R key operation

The entire recorder operates under the control of a standard 80 C31 microprocessor 30 which, besides

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its internal memory is provided with external memory in the form of a 32 kb EPROM 32 and an 8 kb RAM 34. A reset circuit 36, to which is connected the ON/OFF key 19, functions to provide a proper startup reset for the system, to block an accidental reset during normal system operation, and to generate an interrupt instead of a reset to shut the system down via the ON/OFF key 10. The beeper circuit 38 is a simple 4.6 KHz beeper which provides audible alarm signals under the control of the microprocessor 30, while the serial interface 40 is the standard serial interface and is used to interface the system with the interface unit.

All the other components of the system are connected to, and controlled by, the microprocessor 30 by a two wire serial bus 42. The EEPROM memory 44 is divided into two parts, in one of which is stored the patient therapy information such as insulin injection times and amounts, blood glucose measurement times, diet meal times and amounts, exercise times and duration. The other part of the EEPROM is used as a memory buffer to store the results of the last 85 blood glucose measurements, thus allowing the patient to review his blood-sugar count over a period of time. As this memory is an EEPROM, the stored data is not lost in the event of a total power loss.

The clock circuit is a PCF8573 integrated circuit suitable for serving the bus 42. The keyboard 48 comprises the keys shown in Fig. 1 with the exception of the ON/OFF key 19 which, as previously noted, is connected directly to the reset circuit 36. Associated with the keyboard is a keyboard decoder 50 which comprises an integrated circuit able to read and drive the keyboard circuit.

The LCD 54 is shown in detail in Fig. 2 and has a driver circuit 52. The sensor circuit 58 with an associated A/D converter are used for blood glucose measurements and will be described in more detail

The operation of the display and the key board for the various functions is as follows:-

INSULIN THERAPY

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At the programmed time an internal, audible alarm goes off and repeats after one minute.

The patient turns on the recorder. by pressing "ON/OFF" key 19 and the display shows the actual time, the alarm symbol (if selected) and the insulin therapy symbol, the insulin symbol and the dots in time blinking slowly.

When the patient presses the insulin key 11, the insulin symbol in the display stops blinking, the insulin type and number of units appear in the display.

After four seconds, the "+" symbol appears in the alphanumeric field and then, in the case of a mixed shot, the second type and number of units appear in the display.

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INSULIN THERAPY FOLLOW UP

When the patient has taken the correct dose of insulin, he presses the red M-store key 21 to confirm 40

In the display the first insulin type and dose is redisplayed followed by "+" and then the second insulin this fact. type and dose in case of a mixed shot. This scroll is done once. Simultaneously "M" appears in the display, together with insulin symbol.

During memory record, the memory may be cleared by pressing the "C" key 15.

After the therapy has been scrolled once, the actual time shows up in the display and insulin symbol vanishes.

INSULIN THERAPY DEVIATION

If the patient wishes to adjust the amount of insulin, he presses the insulin key 11 and the first insulin type and dose appears in display. The patient can then press the "+" key 17 or the "-" key 18 to increase or decrease the insulin units.

If there is a second type of insulin (i.e. a mixed shot) the insulin key 11 can be pressed again and the dose adjusted in the same way as above.

When the dose has been taken, the red "M" key 21 is pressed to confirm and store the adjusted

therapy. The display confirms "adjusted" therapy by scrolling adjusted therapy once, displaying solid insulin bottle and "M" for memory record. The display then returns to actual time display.

The "C" key 15 can always be used to clear numbers currently being displayed. This mode is identical to normal calculator operation. To return to the prescribed therapy after deviation, patient presses the "R" key 16.

DIET THERAPY

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The diet therapy may be operated in any one of three ways:-

- A. Diet exchange system: individually selected exchanges per meal.
- B. Diet indication system requiring an entry indicating:

above normal diet "+" key 17

normal diet "R" key

below normal diet "-" key 18

C. Diet "YES/NO" system requiring an entry indicating :

diet followed "M" key 21

diet skipped "C" key 15.

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PROCEDURE FOR DIET EXCHANGE

The procedure is essentially the same as for the insulin therapy. At the programmed time the audible alarm sounds and is repeated after one minute. The patient turns on the monitor/recorder by pressing the "ON/OFF" key and the display shows the actual time, the alarm signal (if selected) and the diet therapy signal. The diet symbol and the dots in time blink slowly.

When the patient presses the diet key 12, the diet symbol stops blinking and the programmed therapy is displayed. In the present invention the programmed diet therapy can be any one of eight possibilities of which six are standard and two can be designed by the doctor.

DIET THERAPY FOLLOW UP

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This is essentially the same procedure as was described for insulin. After completing the terapy the red "M" key 21 is pressed to record the fact, the diet therapy scrolls once while "M" appears in the display with a steady diet symbol. Thereafter the actual time shows in the display.

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DIET THERAPY DEVIATION

The therapy can be adjusted by means of the "+" and "-" keys as described for insulin therapy.

PROCEDURE FOR DIET INDICATION AND DIET YES/NO SYSTEMS

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The procedures for these possible systems (the initial choice being made by the master computer) is essentially the same as described above, the difference being that in the diet indication system the patient must enter whether the food eaten is above, below or equal to the normal programmed diet, and in the diet YES/NO system the patient must indicate whether the diet was followed or skipped.

EXERCISE THERAPY

This may be programmed in any one of three ways.

- A. Specific exercise system with individually selected exercise level and duration, set in Homer and downloaded to Romeo.
 - B. Exercise indication system requiring entry indicating:
- "+" key - more exercise
- "R" key - normal exercise
- "-" key. - less exercise
 - C. Exercise "YES/NO" system requiring entry indicating:
 - "M" key - exercise done
 - exercise skipped "C" key.

PROCEDURE

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The procedure for exercise therapy is essentially the same as for the therapies previously described.

URINE TESTING

The master computer will programme the monitor/recorder in one of four ways, viz: 25

- -if this function is used at all
- -if only ketones function is used
- -if only urine glucose is used
- -if both ketones and urine glucose are used.

Any test strip can be used as the patient has always to make a manual entry for this function. The 30 master computer must be informed of the type of strip(s) being used and the appropriate coding.

PROCEDURE

The procedure followed by the patient is essentially the same as previously described except that the patient must enter the results of the test. In the case of the urine ketone function a negative test result is recorded by use of the "-" key 18 while a positive test result is recorded by use of the "+" key 17 once, twice or three times.

In the case of the urine glucose function, the "-" key 18 is used to record a negative result while the "+" key 17 is used up to eight times to record the possible urine glucose % values as follows: less than 0.1, 0.1 to 0.5, 0.5 to 1.0, 1.0, 1.0 to 2.0, 2.0 to 3.0, 3.0 to 4.0, 4.0 to 5.0, 5.0 to 9.9, and greater than 9.9.

BLOOD GLUCOSE MEASUREMENT

This measurement is effected automatically by a reader in a special compartment in the recorder which will be described in more detail later.

PROCEDURE

The procedure commences with the sounding of the alarm and the normal response by the patient of pressing the "ON/OFF" key 19, the therapy key (blood glucose 10) and inserting a clean test strip into the compartment. The therapy key is again depressed. This allows the recorder to obtain a standard or base

reading. Thereafter the patient using the strip from the compartment proceeds with the test by pricking his finger, blotting it on the test strip, and pressing the blood glucose key 10. The recorder then counts for 60 seconds giving audible warnings for the last few seconds to enable the patient to clean the strip at 60 seconds. The recorder continues to count to 120 seconds and gives audible warnings from 115 seconds to enable the patient to insert the strip into the special compartment.

The subsequent measurement of the blood glucose value and the entry of the measured value into memory is automatic.

10 EVENT MARKERS

An event marker is an additional entry made on the initiative of the patient to record significant events such as

- "+" & BG keys hyper
- "+" & "diet" keys illness
- "+" & "exercise" keys stress
- " + " & "insulin" keys fever
- "+" & "urine" keys alcohol

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RECALL FUNCTION

The recall function enables the patient to go back to an earlier therapy step and store the corresponding activity so long as there is no same therapy alarm active.

RECALL ROUTINE

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The patient turns on the recorder and presses the "R" key 16 and then the desired therapy key. The recorder then goes back one therapy in time to an unrecorded blank therapy step and displays the programmed time of that step. The patient can then either accept that time by pressing the therapy key again or can adjust the time to the time at which the therapy took place.

THE BLOOD GLUCOSE TEST STRIP READER

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The blood glucose test strip reader 60 is illustrated in Fig. 4 and is located in the middle of the bottom edge of the recorder. The reader 60 consists of three superimposed parts, viz. the strip carrier 61 which positions the BG test strip correctly in the light path, the reflection compartment 62 which defines the light compartment, and an emitter/sensor unit which transmits and receives the light.

The emitter/sensor unit comprises a housing 63 in which are located a light emitting diode 64 having a light intensity output at 565 nM and a photodiode 65 sensitive to the green light of that wavelenght. Above each of the diodes 64 and 65 is located an aperture, 66 and 67 respectively, allowing light to pass to and from the reflection compartment, the aperture 66 being formed so that the light from the LED 64 emerges from the housing 63 as a narrow high intensity beam. The housing 63 is accurately located in the main body 68 of the recorder so that the diodes are accurately located relative to the reflection compartment and the strip carrier.

The reflection compartment (see Figs. 4 and 7) comprises a calibrated slit 70 immediately above the housing 63, a glass 71 which has an anti-reflection coating with optimum light transmission at 565 nM, and a strip support 72 having a slit 73 above the calibration slit 70. The support 72 serves to space the bloody test strip from the glass 71.

With this arrangement of the reflection compartment above the emitter/sensor unit it can be seen that a strip, accurately located on the support 72 above the slit 73 has fixed spatial relationship with the diodes 64 and 65. It is the purpose of the strip carrier to ensure accurate location of the strip.

The strip carrier 61 as seen in Fig. 5 is essentially a square cover provided with two positioning tongues 75 at its inner edge, the tongues being tapered in two directions to cooperate with two similar cavities 76 in the main body 68 of the recorder thereby to locate accurately the inner edge of the carrier in the recorder. The outer edge of the carrier is provided with clamps 77 which have upper surfaces 78 spaced below the main part of the carrier. The carrier 61 is slid horizontally into position, guide rails (not illustrated) on the main body 68 of the recorder engaging its lateral sides. When fully in position the upper surfaces 78 of the clamps engage the bottom surfaces of the rails thus securely and accurately locating the strip carrier 61 in the recorder body 68.

The strip carrier 61 is provided with a slit 80 whose width is the same as that of a blood glucose test strip. A cantilevered tongue 81 overlies the main part of the slit 80, extending outwardly from the inner end wall 82 of the slit 80. The tongue 81 is thinner than the main body of the carrier so that a gap 85, substantially equal to the thickness of a test strip, is provided between the bottom of the tongue 81 and the upper surface of the strip carrier 71. To assist in locating a test strip in the gap 81 the side walls of ite slit 80 have inclined lead-in portions 86 and the underside of the tongue 81 is formed as a lead-in ramp 87.

When the test strip reader is assembled as shown in Fig. 4, a test strip is inserted into the slit 80 beneath the tongue 81 and when the end of the strip abuts the end wall 82 of the slit 80 the active portion of the strip is located above the diode 64 to be illuminated thereby, light reflected from the active portion being detected by the diode 65.

As seen in Fig. 8 the light emitting diode 64 is energised from a D/A - A/D converter 90, the output from the photodiode 65 passing to an operational amplifier whose output is connected back to the converter 90.

OTHER FUNCTIONS

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The recorder also functions to remind a patient that a consultation with the doctor is due by displaying the telephone symbol, that the memory is getting full by displaying the memory full up symbols, and that the battery is becoming exhausted by displaying the battery low symbol.

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SOFTWARE

The recorder operates under the supervision of a terminal software programme stored in the 32K EPROM and the recorder's operation is determined by a software profile downloaded into the EEPROM.

The terminal software consists of the following sections:-

- a) blood glucose test strip sensor response conversion
- b) microprocessor soft control
- c) LCD soft driver
- d) keyboard encoder/decoder
 - e) data compression
 - f) diagnostics
 - g) power control

The software profile consists of the following sections:-

- a) key function : defines if the keys are operational or not
- b) key reaction : defines what action follows activation of a particular key
- c) guidelines : defines alarm times, insulin doses and types diet specifications and exercise
- d) sensor setting : defines which blood glucose test strip can be used in the corresponding response specifications curve. 50

The data compression techniques employed are designed to save storage space by a coding system which uses parts of a single byte to refer to different aspects of a therapy. For example in insulin therapy the following coding is used:-

* DIAMETIC DATA STORAGE —INSULIN— *

5	-	λ	PRESCRIPTION FOLLOW UP		
10			00177700	00 XXX 001	= INSULIN CODE = SHOT NUMBER = FOILOW UP CODE
	-	В	TIMING DEVIATION		
			00177701	001 YYY 01	= INSULIN CODE = SHOT NUMBER = TIMING DEVIATION CODE
15			********	WANAAA	= 0-239 UNITS OF 6 MIN.
	-	c	TIMING AND THERAPY DEVIATION		
20			00144410	001 YYY 10	= INSULIN CODE = SHOT NUMBER = TIMING & THERAPY DEVIATON CODE
25			HAMPANAM DODDDDDD f or f or m DODDDDDD m or l or l	DECCESSOR	= 0-239 UNITS OF 6 MIN. = INSULIN DOSE 0-253 * 0.5 UNITS
	-	D	ADDITIONAL ENTRY		
30			00100011	001 00011 MANAGEM DDDDDDDDD	= INSULIN CODE = ADDITIONAL ENTRY CODE
			MINIMAN DDDDDDDDD f or f or m DDDDDDDDD m or 1 or 1		= 0-239 UNITS OF 6 MIN. = INSULIN DOSE 0-253 * 0.5 UNITS

A similar pattern is used for the other therapies available - the first three bits of the byte defining the therapy, the second three defining the therapy number and the last two indicating the number of relevant bytes to follow.

A full programme listing for the recorder is included in Appendix A hereto, and the software documentation is included in Appendix B.

INTERFACE

Introduction

As previously discussed the recorder is used with an interface unit to enable data to be transferred between the recorder and the master computer. The recorder simply plugs into the interface which in turn plugs into the master computer or a modem. Alternatively the data can be printed out on a printer which can be incorporated in the interface.

Referring to Fig. 9, the interface comprises the following components.

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Blocks 100, 102 and 103

The blocks contain the common known microcontroller components such as a microprocessor (100), address-latches (102) and a static ram (103).

Block 104

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When the interface is switched on, this module provides a reset for the microprocessor in order to read out the program-memory from the proper start address.

Block 105

With the programmable Peripheral Interface (P.P.I.) it is possible to connect a thermal printer to print out diabetic data. This integrated circuit has three parallel I/O ports. Two are used for the optional printer. The third part is used for indication of the present Juliet-mode for which LED's are used. Every time only one LED lights up next to a symbol to indicate this mode.

Block 106 The programmable Interrupt controller (P.I.C.)

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Handles several interrupts generated by other peripherals such as the keyboard or the serialcommunication port which is desribed in Block 107. On its turn the P.I.C. generates an interrupt which is sent to the microprocessor.

If the microprocessor receives an interrupt the controller is checked for the origin.

Block 107

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The Programmable Communications Interface (P.C.I.) provides together with Block 108 the serial communication with a computer. The P.C.I. can be programmed to communicate at several baud-rates.

For direct communication with a computer 9600 baud is used.

It is possible to connect a modern to this port in order to communicate over long distances. The transmission speed is selectable: 1200 baud or 300 baud.

Block 108

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This block contains the drivers and receivers to transform T.T.L. to V.24 levels and vice versa.

Block 109

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The key-board has three push-buttons. They have special symbols according to their function.

- : Scrolling through the several modes.
- : Actual start of the operation.
- : Paper feed (with printer option). 55

Block 110

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The mode indications of Juliet are listed below. Print-out of blood-glucose values in graphic and in list format. Print-out of insulin values in graphic and in list format. Print-out of diet exchange values in graphic and in list format. Print-out of exercise values in graphic and in list format. Print-out of summary of the last seven days recording. Therapy listing. 10 Phone-modem communication at a speed of 1200 baud. Phone-modem communication at a speed of 300 baud. Direct communication with a computer at a speed of 9600 baud.

Block 111

An optional printer-box can be inserted. For the possible print-outs see above.

Block 112

The interface is powered by an external transformer (220/110 V to 2 x 6 V). 25 Within the interface this voltage is stabilized to the several necessary voltage-levels.

Block 113

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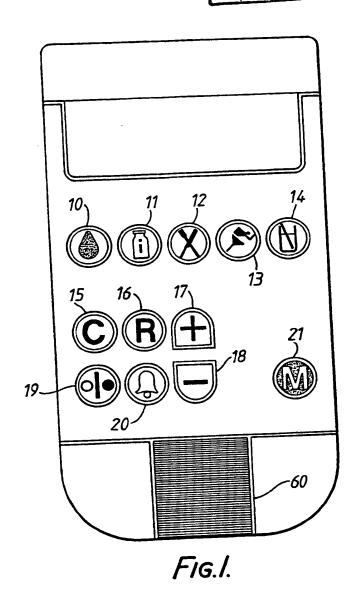
The Romeo-Juliet communication is realized by a 6 pins connector, which is connected to the serial port of the microprocessor.

A full programme listing for the interface is provided in Appendix C.

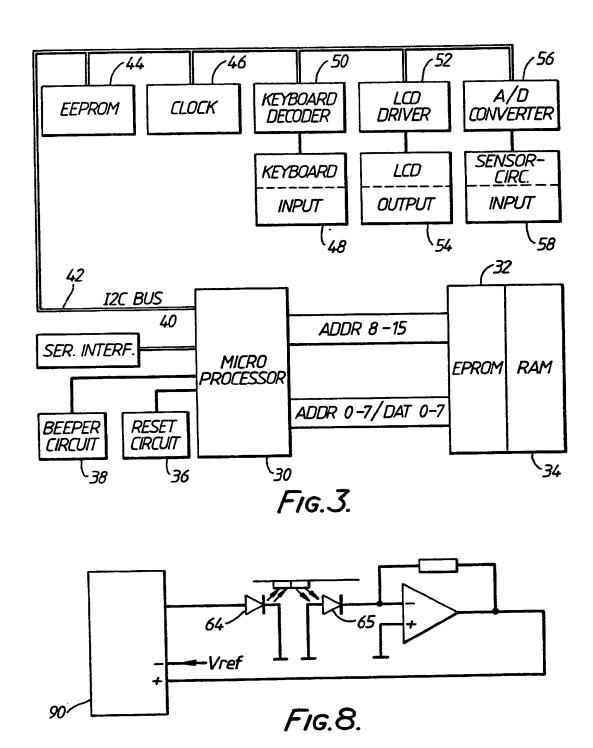
Claims

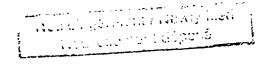
- 1. A system of outpatient management comprising a computer into which are loaded details of the patient, or his condition, a monitor in the possession of the patient and an interface, enabling the computer and the monitor to exchange information, the computer being programmed to produce from the loaded patient details a course of treatment or medication which is recorded in the patient's monitor, the monitor including a computer, a display and a keyboard and being programmed to use the recorded course to display at appropriate times the desired treatment or medication and to accept and record information entered by the patient on the keyboard.
 - 2. A monitor for use by a patient in a system of outpatient management, the monitor comprising a computer, a keyboard and a display, the computer being being adapted to store a course of medication of treatment to display at appropriate times the desired medication or treatment and to accept and store keyboard input from the patient relating to the medication or treatment.
 - 3. Apparatus for measuring colour change comprising means for supporting an article subject to colour change, a light source and means for directing light from the source onto the article as a well defined spot, and means for receiving reflected light from the light source and measuring the intensity of the light.
 - 4. Apparatus as claimed in claim 3, in which the article is a blood glucose test strip.
- 5. Apparatus as claimed in claim 3 or claim 4, including an optical filter located between the article and the light source and having optimum transmission at the dominant wavelength of the source.

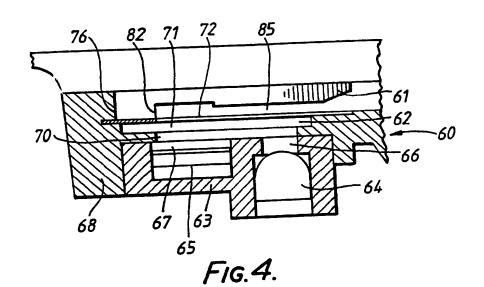
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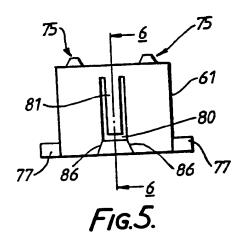


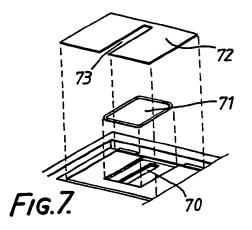
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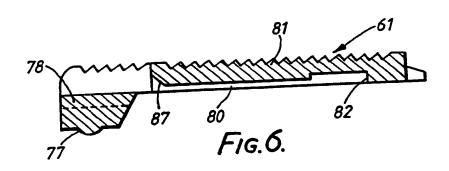


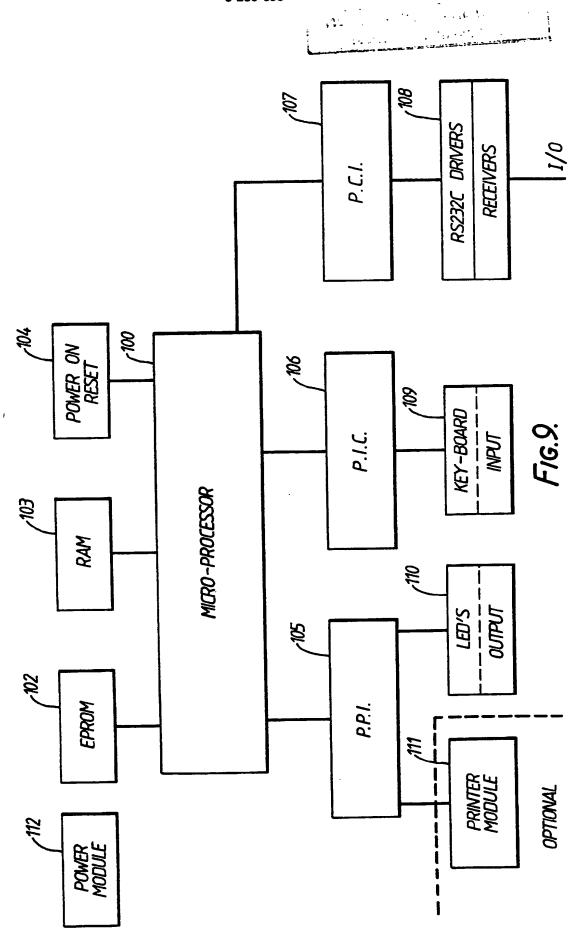












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- Diabetes management system and apparatus.
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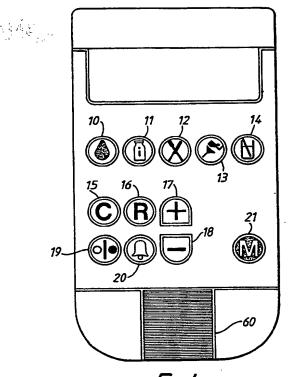


FIG.I.

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EUROPEAN SEARCH REPORT

EP 87 30 4029.

	DOCUMENTS CONSID	dication, where approp	clevani	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. CI.4)	
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	* Page 1, lines 4- 9 - page 4, line 2-14; page 11, 12, lines 22-35	e 3; page 5 Lines 14-35	, lines	1,2		
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	Place of search		ion of the search	'	LECOMTE	
<u> </u>	The Hague			r principle ur	dedving the invention	
CATEGORY OF CITED DOCUMENTS X: particularly relevant if taken alone Y: particularly relevant if combined with another document of the same category A: technological background O: non-written disclosure P: intermediate document			T: theory or principle underlying the invention E: earlier patent document, but published on, or after the filing date D: document cited in the application L: document cited for other reasons			
A	technological background non-written disclosure intermediate document		&: member	of the same	patent family, corresponding	



CL	AIMS INCURRING FEES
The presen	t European patent application comprised at the time of filing more than ten claims.
	All claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for all claims.
	Only part of the claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims and for those claims for which claims fees have been paid.
	namely claims:
	No claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims.
	·
X LA	CK OF UNITY OF INVENTION
	n Division considers that the present European patent application does not comply with the requirement of unity of and relates to several inventions or groups of inventions,
1. c	laims 1,2: Monitor for outpatient management
2. C	laims 3-5: Apparatus for measuring color change
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	All further search fees have been paid within the fixed time limit. The present European search report has been drawn up for all claims.
	Only part of the further search fees have been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the inventions in respect of which search fees have been paid,
	namely claims:
四	None of the further search fees has been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the invention first mentioned in the claims,
	namely claims: 1,2

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